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FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 11:21:33 ON 28  
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L1 20802 S LYSOPHOSPHATIDYLCHOLINE  
L2 1046 S L1 AND ATHEROSCLEROSIS?  
L3 0 S L1 AND HYPERTWNSION?  
L4 39 S L2 AND HYPERTENSION?  
L5 1152 S L1 AND (CARDIOVASCULAR?)  
L6 304 S L2 AND L5  
L7 921 S L1 AND PHOSPHOCHOLINE?  
L8 2 S L6 AND L7  
L9 2 DUPLICATE REMOVE L8 (0 DUPLICATES REMOVED)  
L10 9 S L5 AND L7  
L11 9 DUPLICATE REMOVE L10 (0 DUPLICATES REMOVED)  
L12 7 S L11 NOT L9  
L13 0 S L7 AND (METABOLIC SYNDROME)  
L14 0 S L13 AND HEART?  
L15 43 S L7 AND HEART?  
L16 25 DUPLICATE REMOVE L15 (18 DUPLICATES REMOVED)  
L17 17 S L16 AND PD<1999

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d his

(FILE 'HOME' ENTERED AT 11:17:00 ON 28 FEB 2007)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 11:21:33 ON 28 FEB 2007

|     |         |  |
|-----|---------|--|
| L1  | 20802 S | LYSOPHOSPHATIDYLCHOLINE                      |
| L2  | 1046 S  | L1 AND ATHEROSCLEROSIS?                      |
| L3  | 0 S     | L1 AND HYPERTWNSION?                         |
| L4  | 39 S    | L2 AND HYPERTENSION?                         |
| L5  | 1152 S  | L1 AND (CARDIOVASCULAR?)                     |
| L6  | 304 S   | L2 AND L5                                    |
| L7  | 921 S   | L1 AND PHOSPHOCHOLINE?                       |
| L8  | 2 S     | L6 AND L7                                    |
| L9  | 2       | DUPLICATE REMOVE L8 (0 DUPLICATES REMOVED)   |
| L10 | 9 S     | L5 AND L7                                    |
| L11 | 9       | DUPLICATE REMOVE L10 (0 DUPLICATES REMOVED)  |
| L12 | 7 S     | L11 NOT L9                                   |
| L13 | 0 S     | L7 AND (METABOLIC SYNDROME)                  |
| L14 | 0 S     | L13 AND HEART?                               |
| L15 | 43 S    | L7 AND HEART?                                |
| L16 | 25      | DUPLICATE REMOVE L15 (18 DUPLICATES REMOVED) |
| L17 | 17 S    | L16 AND PD<1999                              |

=>

ANSWER 14 OF 17 MEDLINE on STN

AN 89322894 MEDLINE

DN PubMed ID: 2665794

TI Regulation of phosphatidylcholine metabolism in mammalian hearts

AU Hatch G M; O K; Choy P C

CS Department of Biochemistry, Faculty of Medicine, University of Manitoba, Winnipeg, Canada.

SO Biochemistry and cell biology = Biochimie et biologie cellulaire, (1989 Feb-Mar) Vol. 67, No. 2-3, pp. 67-77. Ref: 104

Journal code: 8606068. ISSN: 0829-8211.

CY Canada

DT Journal; Article; (JOURNAL ARTICLE)  
(RESEARCH SUPPORT, NON-U.S. GOV'T)  
General Review; (REVIEW)

LA English

FS Priority Journals

EM 198908

ED Entered STN: 9 Mar 1990  
Last Updated on STN: 9 Mar 1990  
Entered Medline: 31 Aug 1989

AB Phosphatidylcholine is the major phospholipid in the mammalian heart. Over 90% of the cardiac phosphatidylcholine is synthesized via the CDP-choline pathway. The rate-limiting step of this pathway is catalyzed by CTP:phosphocholine cytidyltransferase. Current evidence suggests that phosphatidylcholine biosynthesis in the heart is regulated by the availability of CTP and the modulation of cytidyltransferase activity. Phosphatidylcholine is degraded mainly by the actions of phospholipase A1 and A2, with the formation of lysophosphatidylcholine. Lysophosphatidylcholine may be further deacylated by lysophospholipase or reacylated back into the parent phospholipid by the action of acyltransferase. The accumulation of lysophosphatidylcholine in the heart may be one of the biochemical factors for the production of cardiac arrhythmias.

CT Animals  
\*Heart: PH, physiology  
\*Mammals: ME, metabolism  
Mammals: PH, physiology  
\*Myocardium: ME, metabolism  
\*Phosphatidylcholines: ME, metabolism  
Phosphatidylcholines: PH, physiology

CN 0 (Phosphatidylcholines)



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 Cover title: Canadian journal of biochemistry and cell biology =

**Author:** Canadian Biochemical Society.  
 National Research Council Canada.  
 Canadian Society for Cell Biology.

**Imprint:** Ottawa : National Research Council of Canada = Conseil national des recherches du Canada, 1986-

**URL:** <http://search.epnet.com/direct.asp?jid=35G&db=aph> Click here for Online version via Academic Search Premier (ASP). Feb 2001-  
<http://proquest.umi.com/pqdweb?RQT=318&VName=PQD&clientid=19649&pmid=36120> Click here for Online version via Proquest. Jan 1, 1998-Present.

**Notes:** Available on ADONIS, v. 73, no. 1-2 (1995) - v. 80, no. 4 (2002)  
 Includes bibliographies.  
 Articles in English; summaries in English and French.  
 Official journal of the Canadian Biochemical Society and the Canadian Society for Cell Biology.

**ISSN:** 0829-8211

**Subjects:** Biological chemistry -- Periodicals.  
 Cytology -- Periodicals.

**Description:** v. : ill. ; 26 cm.

**Continues:** Canadian journal of biochemistry and cell biology

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ANSWER 1 OF 1 MEDLINE on STN

AN 92197126 MEDLINE

DN PubMed ID: 1801455

TI [Phospholipid thrombocyte activating factor, its analogs and antagonists: prospects of their use in medicine].  
Fosfolipidnyi faktor aktivatsii trombotsitov, ego analogi i antagonisty: perspektivy primeneniia v meditsine.

AU Kulikov V I; Muzia G I

SO Vestnik Akademii meditsinskikh nauk SSSR, (1991) No. 10, pp. 13-7. Ref: 37

Journal code: 7506153. ISSN: 0002-3027.

CY USSR

DT (ENGLISH ABSTRACT)  
(IN VITRO)  
Journal; Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)

LA Russian

FS Priority Journals

EM 199204

ED Entered STN: 9 May 1992  
Last Updated on STN: 9 May 1992  
Entered Medline: 21 Apr 1992

AB Experimental data on the biological activity of phospholipid platelet-activation factor (PAF), its structural analogs and antagonists are discussed. The prospects of the use of PAF and PAF antagonists in medicine are under consideration. The conclusion is drawn that PAF antagonists may serve the basis for the development of highly potent drugs of new generation.

CT Azepines: DU, diagnostic use  
Azepines: PD, pharmacology  
\*Azepines: TU, therapeutic use  
\*Diterpenes  
Fibrinolytic Agents: PD, pharmacology  
\*Fibrinolytic Agents: TU, therapeutic use  
Ginkgolides  
Humans  
Lactones: DU, diagnostic use  
Lactones: PD, pharmacology  
\*Lactones: TU, therapeutic use  
\*Lysophosphatidylcholines: PD, pharmacology  
\*Platelet Activating Factor: AA, analogs & derivatives  
Platelet Activating Factor: AI, antagonists & inhibitors  
\*Platelet Activating Factor: PH, physiology  
Platelet Activation: DE, drug effects  
\*Platelet Activation: PH, physiology  
Platelet Aggregation: DE, drug effects  
\*Platelet Aggregation: PH, physiology  
Platelet Function Tests  
Thrombosis: BL, blood  
Thrombosis: DT, drug therapy  
\*Thrombosis: ET, etiology  
Triazoles: DU, diagnostic use  
Triazoles: PD, pharmacology  
\*Triazoles: TU, therapeutic use

RN 105219-56-5 (WEB 2086); 99796-69-7 (ginkgolide B)

CN 0 (1-acylglycerolphosphorylcholine); 0 (1-alkyl-2-acyl-sn-glycero-3-phosphocholine); 0 (Azepines); 0 (Diterpenes); 0 (Fibrinolytic Agents); 0 (Ginkgolides); 0 (Lactones); 0 (Lysophosphatidylcholines\*\*  
\* ); 0 (Platelet Activating Factor); 0 (Triazoles)

=>

ANSWER 2 OF 22 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN

AN 1990:514666 BIOSIS

DN PREV199090131942; BA90:131942

TI HYDROLYSIS OF 2 ACYL-SN-GLYCERO-3-PHOSPHOCHOLINES IN GUINEA-PIG  
HEART MITOCHONDRIA.

AU BADIANI K [Reprint author]; PAGE L; ARTHUR G

CS DEP BIOCHEM MOL BIOL, FAC MED, UNIV MANITOBA, 770 BANNATYNE AVE, MANIT,  
CANADA R3E 0W3

SO Biochemistry and Cell Biology, (1990) Vol. 68, No. 9, pp.  
1090-1095.  
CODEN: BCBIEQ. ISSN: 0829-8211.

DT Article

FS BA

LA ENGLISH

ED Entered STN: 19 Nov 1990  
Last Updated on STN: 19 Nov 1990

AB Although both 2-acyl-sn-glycero-3-phosphocholine and  
1-acyl-sn-glycero-3-phosphocholine may be produced from  
phosphatidylcholine hydrolysis, studies on the former have lagged behind  
that of the latter. In this study a lysophospholipase A2 that hydrolyses  
2-acyl-sn-glycero-3-phosphocholine has been characterized in  
guinea pig heart mitochondria. The lysophospholipase A2 activity was not  
dependent on Ca<sup>2+</sup> and was inhibited differentially by saturated and  
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discriminate among different molecular species of 2-acyl-sn-glycero-3-  
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The order of decreasing rates of hydrolysis of different molecular species  
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presented singly, was 18:2 > 20:4 > 18:1 > 16:0. A differential  
inhibition of the rate of hydrolysis of the individual substrates was  
observed when the substrates were presented in pairs. The degree of  
inhibition was dependent on the molar ratio of the mixed substrates. The  
characteristics of the enzyme suggest that involvement in the selective  
release of fatty acids from mitochondrial phosphatidylcholine would depend  
on a high selectivity of phospholipase A1 for different molecular species  
of phosphatidylcholine. A lysophospholipase A1 activity was also  
characterized in the mitochondria with a distinct acyl specificity from  
the lysophospholipase A2. Other characteristics of the two  
lysophospholipases suggest that the two reactions are not catalyzed by the  
same enzyme.

CC Biochemistry studies - Proteins, peptides and amino acids 10064  
Biochemistry studies - Lipids 10066  
Enzymes - Physiological studies 10808  
Anatomy and Histology - Microscopic and ultramicroscopic anatomy 11108  
Metabolism - Lipids 13006  
Cardiovascular system - Physiology and biochemistry 14504

IT Major Concepts  
Cardiovascular System (Transport and Circulation); Enzymology  
(Biochemistry and Molecular Biophysics); Metabolism; Morphology

IT Miscellaneous Descriptors  
FATTY ACID RELEASE

ORGN Classifier  
Caviidae 86300  
Super Taxa  
Rodentia; Mammalia; Vertebrata; Chordata; Animalia  
Taxa Notes  
Animals, Chordates, Mammals, Nonhuman Vertebrates, Nonhuman Mammals,  
Rodents, Vertebrates

ANSWER 2 OF 22 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN

AN 1990:514666 BIOSIS

DN PREV199090131942; BA90:131942

TI HYDROLYSIS OF 2 ACYL-SN-GLYCERO-3-PHOSPHOCHOLINES IN GUINEA-PIG  
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Metabolism - Lipids 13006  
Cardiovascular system - Physiology and biochemistry 14504

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IT Miscellaneous Descriptors  
FATTY ACID RELEASE

ORGN Classifier  
Caviidae 86300  
Super Taxa  
Rodentia; Mammalia; Vertebrata; Chordata; Animalia  
Taxa Notes  
Animals, Chordates, Mammals, Nonhuman Vertebrates, Nonhuman Mammals,  
Rodents, Vertebrates




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**Author:** Canadian Biochemical Society.  
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